Claims

1. A pharmaceutical composition comprising (i) a first specific binding agent selected from an antibody or a large binding fragment of an antibody which specifically binds a target toxin, and (ii) a second specific binding agent which comprises a small binding fragment of an antibody which binds said toxin.

2. A composition according to claim 1 wherein the first specific binding agent comprises a large binding fragment of an antibody.

3. A composition according to claim 2 wherein the large binding fragment of an antibody is a $F(ab')_2$ or $F(ab)_2$ fragment.

4. A composition according to claim 1 wherein the first specific binding agent is an antibody which is an IgG or IgT.

5. A composition according to claim 4 wherein the antibody is humanised.

6. A composition according to any one of the preceding claims wherein the second specific binding agent comprises an Fab, Fab', a single chain (sc) antibody or an FV, VH or VK fragment.

7. A composition according to claim 6 wherein the second specific binding agent comprises Fab or Fab' fragment.

8. A composition according to any one of the preceding claims wherein the first and/or second binding agents are derived from polyclonal antibodies.

9. A composition according to any one of claims 1 to 7 wherein the first and/or second binding agents are derived from monoclonal antibodies.

10. A composition according to any one of the preceding claims wherein at least one of the first or second specific binding agents includes a section corresponding to part of the Fc region of an antibody.

11. A composition according to any one of the preceding claims wherein the toxin is a Botulinum *toxin*.

12. A composition according to claim 11 wherein the first and second specific binding agents bind at least one of type A, B, C, D, E, F or G botulinum toxin.

13. A composition according to claim 12 wherein the composition comprises sets of first and second specific binding agents each set of specific binding agents binding a different one of botulinum toxins A, B, C, D, E, F or G.

14. A composition according to any one of the preceding claims wherein the w/w ratio of the first specific binding agent to the second specific binding agent is in the range of from 90:10 to 10:90.

15. A composition according claim 14 wherein the w/w ratio of the first specific binding agent to the second specific binding agent is in the range of from 70:30 to 30:70.

16. A composition according to claim 15 wherein the w/w ratio of the first specific binding agent to the second specific binding agent is in the range of from 60:40 to 40:60.

17. A composition according to any one of the preceding claims which further comprises a pharmaceutically acceptable carrier or excipient.

18. A composition according to any one of the preceding claims which is suitable for oral, parenteral, or intranasal

administration, or for administration by inhalation or insufflation.

19. A combination of (i) a first specific binding agent selected from an antibody or a large binding fragment of an antibody which specifically binds a target toxin, and (ii) a second specific binding agent which comprises a small binding fragment of an antibody which binds said toxin, for use in the treatment of the effects of the toxin.

20. The use of a combination of (i) a first specific binding agent selected from an antibody or a large binding fragment of an antibody which specifically binds a target toxin, and (ii) a second specific binding agent which comprises a small binding fragment of an antibody which binds said toxin, in the preparation of a medicament for the treatment of the effects of the toxin.

21. A method of preventing the effects of a toxin on a mammal such as a human, said method comprising administering to a mammal in need thereof, a composition according to any one of claims 1 to 18.

22. A composition substantially as hereinbefore described.